This report is required by law (7 USC 2143). Failure to report according to the regulations can result in an order to cease and desist and to be subject to penalties as provided for in Section 2150.

See reverse side for additional information.

Interagency Report Control No 0180-DOA-AN

UNITED STATES DEPARTMENT OF AGRICULTURE
ANIMAL AND PLANT HEALTH INSPECTION SERVICE

ANNUAL REPORT OF RESEARCH FACILITY

(TYPE OR PRINT)

1. REGISTRATION NO. 50-R-0003 **CUSTOMER NO.** 27

FORM APPROVED
OMB NO. 0579-0036

2. HEADQUARTERS RESEARCH FACILITY *(Name and Address, as registered with USDA, include Zip Code)*

INTERVET INC
29160 INTERVET LANE
MILLSBORO, DE 19966
(b)(2)High, (b)(7)f

3. REPORTING FACILITY (List all locations where animals were housed or used in actual research, testing, teaching, or experimentation, or held for these purposes. Attach additional sheets if necessary.)

FACILITY LOCATIONS*(sites)*

See Attached Listing

RECEIVED
NOV 20 2006

REPORT OF ANIMALS USED BY OR UNDER CONTROL OF RESEARCH FACILITY *(Attach additional sheets if necessary or use APHIS FORM 7023A)*

A. Animals Covered By The Animal Welfare Regulations	B. Number of animals being bred, conditioned, or held for use in teaching, testing, experiments, research, or surgery but not yet used for such purposes.	C. Number of animals upon which teaching, research, experiments, or tests were conducted involving no pain, distress, or use of pain-relieving drugs.	D. Number of animals upon which experiments, teaching, research, surgery, or tests were conducted involving accompanying pain or distress to the animals and for which appropriate anesthetic, analgesic, or tranquilizing drugs were used.	E. Number of animals upon which teaching, experiments, research, surgery or tests were conducted involving accompanying pain or distress to the animals and for which the use of appropriate anesthetic,analgesic, or tranquilizing drugs would have adversely affected the procedures, results, or interpretation of the teaching, research, experiments, surgery, or tests. *(An explanation of the procedures producing pain or distress in these animals and the reasons such drugs were not used must be attached to this report)*	F. TOTAL NO. OF ANIMALS (Cols. C + D + E)
4. Dogs		167	129	56	352
5. Cats		298			298
6. Guinea Pigs	162	258	2121	717	3096
7. Hamsters			385	140	525
8. Rabbits	16	7	2104	479	2590
9. Non-Human Primates					
10. Sheep					
11. Pigs	259	2069			2069
12. Other Farm Animals					
Cattle	342	2362			2362
13. Other Animals					
Equine	84	338			338

ASSURANCE STATEMENTS

1) Professionally acceptable standards governing the care, treatment, and use of animals, including appropriate use of anesthetic, analgesic, and tranquilizing drugs, prior to, during, and following actual research, teaching, testing, surgery, or experimentation were followed by this research facility.

2) Each principal investigator has considered alternatives to painful procedures.

3) This facility is adhering to the standards and regulations under the Act, and it has required that exceptions to the standards and regulations be specified and explained by the principal investigator and approved by the Institutional Animal Care and Use Committee (IACUC). **A summary of all the exceptions is attached to this annual report.** In addition to identifying the IACUC-approved exceptions, this summary includes a brief explanation of the exceptions, as well as the species and number of animals affected.

4) The attending veterinarian for this research facility has appropriate authority to ensure the provision of adequate veterinary care and to oversee the adequacy of other aspects of animal care and use.

CERTIFICATION BY HEADQUARTERS RESEARCH FACILITY OFFICIAL
(Chief Executive Officer or Legally Responsible Institutional official)

(b)(6), (b)(7)c

DATE SIGNED 11/16/06

- HEADQUARTERS

Column E Explanation

This form is intended as an aid to completing the Column E explanation. It is not an official form and its use is voluntary. Names, addresses, protocols, veterinary care programs, and the like, are not required as part of an explanation. A Column E explanation must be written so as to be understood by lay persons as well as scientist.

1 Registration Number _____*50-R-0003*_____

2 Number _____56_____ of animals used in this study

3 Species (common name) ____dogs_____ of animals used in this study

4 Explain the procedure producing pain and/or distress

All dogs were (b)(4) *with a* (b)(4) *for the basis of determining* (b)(4) *efficacy.*

5 Provide scientific justification why pain and/or distress could not be relieved. State methods or means used to determine that pain and/or distress relief would interfere with test results (For Federally mandated testing, see Item 6 below)

Clinical signs, as observed following (b)(4) *are the basis for comparing* (b)(4) *Therefore animals did not receive treatment for clinical signs. Typical clinical signs observed included* (b)(4) (b)(4)

6 What, if any, federal regulations require this procedure? Cite the agency, the code of Federal Regulations (CFR) title number and the specific section number (e.g. APHIS, 9 CFR, 113.102)

Agency _____*9*CFR _____

Column E Explanation

This form is intended as an aid to completing the Column E explanation. It is not an official form and its use is voluntary. Names, addresses, protocols, veterinary care programs, and the like, are not required as part of an explanation. A Column E explanation must be written so as to be understood by lay persons as well as scientist.

1 Registration Number *50-R-0003*

2 Number *717* of animals used in this study

3 Species (common name) *guinea pig* of animals used in this study

4 Explain the procedure producing pain and/or distress

These animals were used for Codified (b)(4) *for the product release testing of all Intervet* (b)(4) *containing products*

Per (b)(4) *the guinea pigs used for* (b)(4) *with* (b)(4) *and observed for 3 days* (b)(4) *and all* (b)(4) *are recorded. The* (b)(4) *causes* (b)(4) *lesions which cause distress.*

Per (b)(4) *the guinea pigs used for* (b)(4) *are* (b)(4) *with* (b)(4) *and observed for 3 days* (b)(4) *and all* (b)(4) *are recorded. The* (b)(4) *causes* (b)(4) *lesions which cause distress.*

In both instances, per code, (b)(4) *'s the endpoint.*

5 Provide scientific justication why pain and/or distress could not be relieved. State methods or means used to determine that pain and/or distress relief would interfere with test results (For Federally mandated testing, see Item 6 below)

(b)(4) *is the end-point. Any intervention must be pre-approved by USDA/APHIS/VS/CVB as written in the filed Outline of Production or Special Outline. Currently, no intervention criteria are written into these procedures.*

6 What, if any, federal regulations require this procedure? Cite the agency, the code of Federal Regulations (CFR) title number and the specific section number (e.g. APHIS, 9 CFR, 113.102)

Agency *USDA/APHIS/VS/CVB* (b)(4)

Column E Explanation

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1 Registration Number 50-R-0003

2 Number 140 of animals used in this study

3 Species (common name) Hamsters of animals used in this study

4 Explain the procedure producing pain and/or distress

These animals were used for (b)(4) *and were* (b)(4) *with* (b)(4)

Per (b)(4) *the hamsters used for passages are* (b)(4)

Per code, (b)(4) *is the endpoint.*

5 Provide scientific justication why pain and/or distress could not be relieved. State methods or means used to determine that pain and/or distress relief would interfere with test results (For Federally mandated testing, see Item 6 below)

(b)(4) *is the end-point.* (b)(4) *occurs within 28/48 hours following the onset of symptoms. Due to the fact that* (b)(4) *is the required endpoint, there are no procedures available to limit discomfort, distress and pain during the challenge period.*

6 What, if any, federal regulations require this procedure? Cite the agency, the code of Federal Regulations (CFR) title number and the specific section number (e.g. APHIS, 9 CFR, 113.102)

Agency *USDA/APHIS/VS/CVB* (b)(4) (b)(4)

Column E Explanation

This form is intended as an aid to completing the Column E explanation. It is not an official form and its use is voluntary. Names, addresses, protocols, veterinary care programs, and the like, are not required as part of an explanation. A Column E explanation must be written so as to be understood by lay persons as well as scientist.

1 Registration Number *50-R-0003*

2 Number *479* of animals used in this study

3 Species (common name) *rabbits* of animals used in this study

4 Explain the procedure producing pain and/or distress

These animals were used for a proprietary (b)(4) *testing for the product release testing of all Intervet* (b)(4) *containing products*

Per our (b)(4) *the rabbits are used in a* (b)(4) (b)(4) *The rabbits are used for* (b)(4) *are* (b)(4) (b)(4) *with* (b)(4) *and observed for 3 days* (b)(4) (b)(4) *and all* (b)(4) *are recorded. The* (b)(4) *causes* (b)(4) *lesions which cause distress. The validity requirement for this test is* (b)(4)

5 Provide scientific justication why pain and/or distress could not be relieved. State methods or means used to determine that pain and/or distress relief would interfere with test results (For Federally mandated testing, see Item 6 below)

(b)(4) *is the end-point parameter measured with this approved* (b)(4) *test. Any intervention must be pre-approved by USDA/APHIS/VS/CVB as would be written in the filed Outline of Production or Special Outline. Currently, no intervention criteria are written into these procedures.*

6 What, if any, federal regulations require this procedure? Cite the agency, the code of Federal Regulations (CFR) title number and the specific section number (e.g. APHIS, 9 CFR, 113.102)

Agency *USDA/APHIS/VS/CVB* (b)(4) *APHIS approved procedure in Intervet* (b)(4)